PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Article 36 and Rule 70)

REG'D 1 AUG 2005

Applicant's or agent's file referen			IVIPO	PCT			
431.F	FOR FURT	HER ACTION	See Form PCT/IPEA/4	16			
International and in the state of				, ,			
International application No. PCT/US2004/024922	International fill 30.07.2004	Ing date <i>(day/month/year)</i>	Priority date (day/mo	onth/year)			
International Patent Classification	n (IPC) or national classificati	on and IPC					
C07F9/6512, C07F9/653, /	461K31/675						
	Applicant						
GILEAD SCIENCES, INC.	et al.						
This report is the internal Authority under Article 3	ational preliminary examin	ation report, established by applicant according to Article	this International Prelimi	inary Examining			
		Abinomit accounting to Willicia	· 36.	.			
3. This report is also account	sheets, including this cover sheet.						
a. Sent to the applie	ANNUALES, COMPRISING:						
sheets of the	and to the international Bureau) a total of sheets, as follows:						
and/or sheet Administrativ	sheets of the description, claims and/or drawings which have been amended and are the basis of this report and/or sheets containing rectifications authorized by this Authority (see Rule 70.16 and Section 607 of the						
sheets which supersede earlier sheets, but which this Authority considers contain an amendment that goes beyond the disclosure in the international application on filed, as indicated to the contain an amendment that goes							
	,,						
sequence listing Box Relating to S	b. (sent to the International Bureau only) a total of (indicate type and number of electronic carrier(s)), containing a sequence listing and/or tables related thereto, in computer readable form only, as indicated in the Supplemental Box Relating to Sequence Listing (see Section 802 of the Administrative Instructions).						
			•				
4. This report contains indi	cations relating to the follo	owing items:					
☐ Box No. I Basis	of the opinion						
☐ Box No. II Priority							
		th regard to novelty, inventiv					
☑ Box No. IV Lack o	of unity of invention	ar regard to novelty, inventive	e step and industrial app	plicability			
🖾 Box No. V Reaso	and the state of t						
☐ Box No. VI Certair	n documents cited	iations supporting such state	ement				
	n defects in the internation	al application					
	n observations on the inter	rnational application					
	·	national application					
Date of submission of the demand		Date of completion of	this report				
		Date of completion of	nno rebott				
17.03.2005		09.08.2005					
Name and malling address of the I	nternational	Authorized Officer					
preliminary examining authority: European Patent Of	ifice			Stirches Petrone			
D-80298 Munich Tel. +49 89 2399 - 0) Tx: 523656 epmu d	Klein, D					
Fax: +49 89 2399 -	1465	Telephone No. +49 89	2399-	The state of the s			

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No. PCT/US2004/024922

_						
_	Box No. I Basis of the	eport				
1	With regard to the language, this report is based on the international application in the language in which it was filed, unless otherwise indicated under this item.					
	international search	n translations from the original language into the following language, of a translation furnished for the purposes of: n (under Rules 12.3 and 23.1(b)) nternational application (under Rule 12.4) inary examination (under Rules 55.2 and/or 55.3)				
2.	. With regard to the elemen have been furnished to the	ts* of the international application, this report is based on (replacement sheets which receiving Office in response to an invitation under Article 14 are referred to in this nd are not annexed to this report):				
	Description, Pages					
	1-133	as originally filed				
	Claims, Numbers					
	1-24	as originally filed				
	☐ a sequence listing and	or any related table(s) - see Supplemental Box Relating to Sequence Listing				
3.	☐ the description, pag☐ the claims, Nos.☐ the drawings, sheet☐ the sequence listing	s <i>f</i> ios				
4.	☐ This report has been est had not been made, since the Supplemental Box (Rule 70)☐ the description, page ☐ the claims, Nos.☐ the drawings, sheets ☐ the sequence listing	stablished as if (some of) the amendments annexed to this report and listed below hey have been considered to go beyond the disclosure as filed, as indicated in the es.				
	* If item 4 applies,	some or all of these sheets may be marked "superseded."				

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

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_	Bo	x No. III Non-establishment				
_		plicability	от ор	inion with regard to novelty, inventive step and industrial		
1.	The obv	he questions whether the claimed invention appears to be novel, to involve an inventive step (to be non- bylious), or to be industrially applicable have not been examined in respect of:				
			he entire international application,			
		claims Nos. 10				
		because:				
		the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (specify):				
		the description, claims or drawings (indicate particular elements below) or said claims Nos. 10 are so unclear that no meaningful opinion could be formed (specify):				
		see separate sheet	parate sheet			
		the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.				
		no international search report has been established for the said claims Nos.				
		the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:				
		the written form		has not been furnished		
				does not comply with the standard		
		the computer readable form		has not been furnished		
				does not comply with the standard		
the tables related to the nucleotide and/or amino acid sequence listing, if in computer renot comply with the technical requirements provided for in Annex C-bis of the Administ			and/or amino acid sequence listing, if in computer readable form only, do ements provided for in Annex C-bis of the Administrative Instructions.			
ı		See separate sheet for further	detail	s		

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International application No. PCT/US2004/024922

_	Box No. IV Lack of unity of invention						
1.	1. In response to the invitation to restrict or pay additional fees, the applicant has:						
		restricted the claims.					
		\square paid additional fees.					
		paid additional fees unde					
		☐ neither restricted nor paid additional fees.					
2.		This Authority found that the requirement of unity of invention is not complied with and chose, according to Rule 68.1, not to invite the applicant to restrict or pay additional fees.					
3.	This Authority considers that the requirement of unity of invention in accordance with Rules 13.1, 13.2 and 13.3 is						
		complied with.					
	☐ not complied with for the following reasons:						
	see separate sheet						
4.	4. Consequently, this report has been established in respect of the following parts of the international application:						
	×						
		the parts relating to claims No	os				
_	Box No. V Reasoned statement under Article 25/2) with reward Asset 1						
_		licability; citations and expl	anatio	ns support	35(2) with regard to novelty, inventive step or industrial ing such statement		
1.	Statement						
	Nov	elty (N)	Yes:	Claims	2,5,8-9,15,18		
			No:	Claims	1,3,4,6,7,11-14,16,17,19-24		
	Inventive step (IS)		Yes:	Claims	2		
		, , ,	No:	Claims	1,3-9,11-24		
Indi		strial applicability (IA)	Yes:	Claims	1-9,11-18,23-24		
		(, , , , , , , , , , , , , , , , , , ,	No:	Claims	19-22		
2.	Cita	tions and explanations (Rule 7	0.7):				

see separate sheet

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (SEPARATE SHEET)

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Re Item II

Priority:

WO 03/087298 and US 2004/014722D1 which are an intermediate document, are not prior art according to the Chap II PCT proceedings.

Nevertheless, the extensive examination of that document, on the question whether it constitutes prior art or not, will depend essentially on the analysis of the claimed priority rights of the present application and will only be performed in the regional European proceedings to come.

Re Item III

Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The subject-matter of claim 10 is unclear contrary to Art. 6 PCT, the reason being that claim 10, wherein R^3 =H, is dependant on claim 3 wherein R^3 = C_1 - C_8 alkyl. Therefore claim 10 will not be examined.

Re Item IV

Lack of unity of invention

This Authority considers that there are 2 inventions covered by the claims indicated as follows:

I: Claims 1(part)-24(part) consisting in compounds bearing one -PY1(Y2H)2 terminal

group (Formula of claim 1 or 11 wherein M2=0 and Rx =

H).

II: Claims 1(part)-24(part) consisting in compounds bearing a terminal group other

than one -PY1(Y2H)2 (Formula of claim 1 or 11 wherein Rx

is different than H).

The common concept linking both invention I and II are antiviral compounds of the formula depicted in claim 11. However this common concept is already known, see for ex D1 (US-A-4 808 716) Formula V or D2 (US 2003/004345) examples 3 and 6.

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY (SEPARATE SHEET)

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Hence, this Authority considers that the above-mentioned inventions are not so linked as to form a single general inventive concept, as required by Rule 13.1 PCT.

Re Item V

Reasoned statement with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

- D1: US-A-4 808 716 (HOLY ANTONIN ET AL) 28 February 1989 (1989-02-28)
- D2: US 2003/004345 A1 (CASARA PATRICK ET AL) 2 January 2003 (2003-01-02)
- D3: EP-A-0 269 947 (BRISTOL MYERS CO) 8 June 1988 (1988-06-08)
- D4: KIM, CHOUNG UN ET AL: "Synthesis and antiviral activity of (S)-9-[4-hydroxy-3- (phosphonomethoxy)butyl]guanine" JOURNAL OF MEDICINAL CHEMISTRY, 33(6), 1797-800 CODEN: JMCMAR; ISSN: 0022-2623, 1990, XP001204580

1) Novelty of Invention I:

D1 discloses compounds of Formula V wherein R1 can be a methyl or hydroxymethyl group and thus anticipate the subject-matter of claims 1,3,4,11,12,13,14,23,24.

D3 discloses compounds of Formulas 3,14,15 and thus anticipate the subject-matter of claims 1,6,7,11,12,16,17,19-24.

D4 discloses on page 56 compound 40 which anticipate the subject-matter of claims 1,11,12,23,24.

Thus claims 1,3,4,6,7,11-14,16,17,19-24 do not comply with the requirements of Art. 33(1) and 33(2) PCT.

2) Inventive step of Invention I:

Claim 2:

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The compounds of claim 2 wherein R^1 - R^8 are each H and wherein M2 is 0 and Rx is H (by extension Y^1 and Y^2 cannot be both O) are not suggested in the prior art. Therefore, claim 2 complies with the requirements of Art 33(3) PCT.

Claims 5,8,15,18:

Claims 5/15 and 8/18 respectively differ form D1/D4 by the length of the R^3/R^5 side chains, more precisely a hydroxyethyl instead of a hydromymethyl in D1/D4.

Since it is common practice of organic chemists who are concerned with the preparation of new drugs to take some compound known to possess some interesting pharmacological properties as a model to study the effect of making changes in its structure, the man skilled in the art would, in the present case, slightly modify the functional group fixed on R³ or R⁵ from D1/D4 (elongate the side chain by one methylene group), and would therefore arrive, without the presence of inventive skill to the subject-matter of claims 5,8,15,18.

Claim 10:

The subject-matter of claim 9 is not considered inventive for the ethylidene group present in the formula is already disclosed in D2. Thus the subject-matter of claim 9 is a mere combination of features known from the prior art.

3) Industrial application:

For the assessment of the present claims 19-22 on the question whether they are industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.